

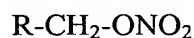
CLAIMS:

1. A method for the treatment or prevention of Alzheimer's disease comprising administering to a subject in need thereof a therapeutically-effective
5 amount of a compound which inhibits the formation or release of β -amyloid and a therapeutically-effective amount of a nitric oxide releaser.

2. The method according to claim 1 wherein the nitric oxide releaser is a nitrate ester of an alkanol.
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3. The method according to claim 1 wherein said inhibitor and said nitric oxide releaser are combined in a single dosage formulation.

4. The method according to claim 3 wherein said inhibitor and said nitric oxide releaser are combined in a compound of formula I:
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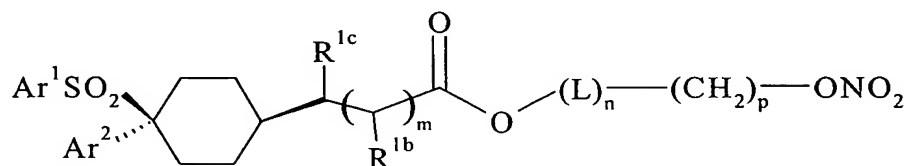


I

where R is such that R-CH₂OH or R-CHO is an inhibitor of the formation or release of A β ;

20 or a pharmaceutically acceptable salt thereof.

5. A compound suitable for use in the method of claim 1, said compound being of formula II:



II

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wherein:

m is 0 or 1;

n is 0 or 1;

L is a linking group;

R^{1c} represents H or C₁₋₄alkyl;

Ar¹ and Ar² independently represent phenyl or heteroaryl, either of which bears 0-3 substituents independently selected from halogen, CN, NO₂, CF₃, CHF₂, OH, OCF₃, CHO, CH=NOH, C₁₋₄alkoxy, C₁₋₄alkoxycarbonyl, C₂₋₆acyl, C₂₋₆alkenyl and C₁₋₄alkyl which optionally bears a substituent selected from halogen, CN, NO₂, CF₃, OH and C₁₋₄alkoxy;

or a pharmaceutically acceptable salt thereof.

6. A compound according to claim 5 wherein m is 1 and R^{1b} and R^{1c} are both H.

15 7. A compound according to claim 5 wherein Ar¹ is 4-chlorophenyl or 4-trifluoromethylphenyl and Ar² is 2,5-difluorophenyl.

8. A compound according to claim 5 wherein L is represented by the formula:

20 -L¹-C(O)O-

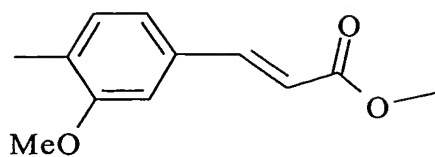
wherein L¹ is a hydrocarbon residue of up to 10 carbon atoms, optionally bearing up to 3 substituents selected from halogen, CN, OH and C₁₋₄alkoxy.

9. A compound according to claim 8 wherein L¹ represents

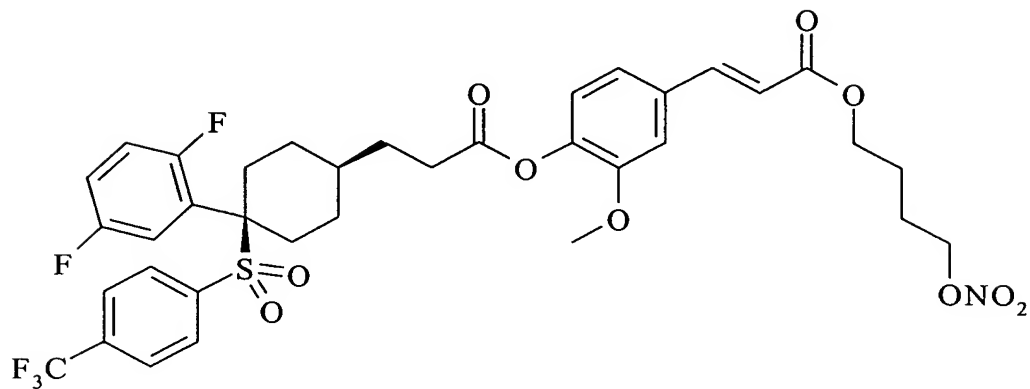
25 $-\text{Ar}-\text{CH}=\text{CH}-$

wherein Ar is a phenyl group bearing up to 2 substituents selected from hydroxy and methoxy.

10. A compound according to claim 5 wherein L represents:

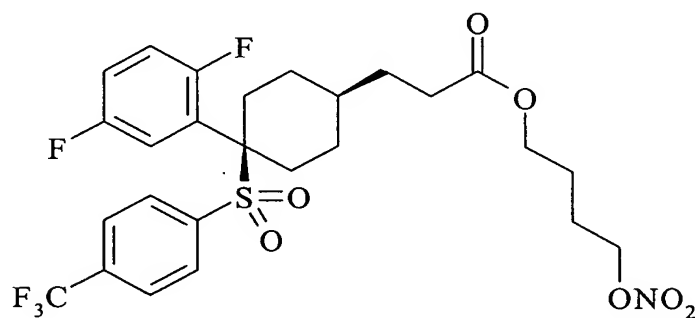


11. A compound according to claim 5 selected from:



5

and



and pharmaceutically acceptable salts thereof.

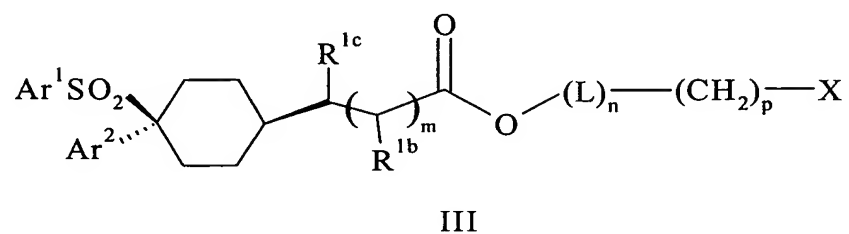
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12. A pharmaceutical composition comprising a compound according to claim 5 and a pharmaceutically acceptable carrier.

13. A method of treatment of a subject suffering from or prone to Alzheimer's disease which comprises administering to that subject an effective amount of a compound according to claim 5.

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14. A method of preparing a compound according to claim 5 comprising reaction of a compound of formula III with silver nitrate:



where X represents chlorine, bromine or iodine, and m, n, p, L, R^{1b}, R^{1c}, Ar¹ and Ar² are as defined in claim 5.